The listing of claims will replace all prior versions and listing of claims in the application:

Listing of Claims:

Claim 1/(currently amended):

A compound, or enantiomers,

stereoisomers and tautomers thereof, or pharmaceutically acceptable salts or solvates of said compound, with said compound having the general structure shown in Formula I:

$$R^{1}$$
 R^{3}
 R^{3}
 R^{2}
 R^{4}
 R^{4

Formula I

wherein M is a moiety of the formula:

having a general structure shown in Formula II:

where k = 0 or 1, and n = 0.5 [[, and p = q = 0, 1 or 2]];

V is a moiety selected from the group consisting of C₁-C₈ alkyl;

 $-(CH_2)_x$ -A- $(CH_2)_y$ -; and $-(CH_2)_c$ -A- $(CH_2)_m$ -C(O)-N(R⁷)- $(CH_2)_d$ -, where A is -O-, -S(O)_r-, and -NR⁷-;

m = 0, 1, 2 or 3; x is a whole number in the range 2-8; y is a whole number in the range 1-5; c is a whole number in the range 2-4; and r = 0, 1 or 2; d is a number in the range 0-5;

[[X and Y are independently selected from the group consisting of N, and CH;]]

one of X is N and the other is CH;

[[Z and Z¹ can be the same or different, each being independently selected from the group consisting of N, CH and N(O);]]

R¹ and R² may each number 1-4 and are independently selected from the group consisting of hydrogen, lower alkyl, lower alkoxy, halogen, polyhalolower alkyl, polyhalolower alkoxy, -OH, CN, NO₂, or COOR⁸; R³ is selected from hydrogen, lower alkyl, lower alkoxy, hydroxyl, with the proviso that when n and k are both 0, then R³ is not -OH or alkoxy;

R⁴ is selected from the group consisting of hydrogen, lower alkyl, polyhalolower alkyl or -OH; and

R⁷ and R⁸ are independently selected from hydrogen, lower alkyl, substituted or unsubstituted phenyl; and substituted or unsubstituted benzyl, wherein said term "substituted" means optional substitution from one or more moieties selected from the group consisting of alkyl, alkoxy, -CF₃, halogen or aryl.

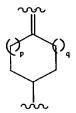
Claim 2 (original): The compound of claim 1, wherein R⁴ is H.

Claim 3 (original): The compound of claim 2, wherein R^1 and R^2 are independently selected from H, halogen, or polyhalolower alkyl.

Claim 4 (canceled).

Claim 5 (amended): The compound of claim 1, wherein M is a

piperazine. [[:



and p = q = 1.]]

Claim & (original): The compound of claim A, wherein \mathbb{R}^4 is H; $\mathbb{R}^1 = \mathbb{R}^2 = \mathbb{H}$, halogen, hydroxy or alkoxy; and R³ is H or lower alkyl.

Claim (original): The compound of Claim 6, wherein $V = C_1 - C_8$ alkyl. Claim \mathcal{L} (original): The compound of claim 5, wherein \mathbb{R}^4 is H; and $\mathbb{R}^1 = \mathbb{R}^2 =$ H, halogen, hydroxy or alkoxy.

The compound of Claim 8, wherein V is $C_1 - C_8$ alkyl. Claim & (original): Claim 12 (previously amended): A pharmaceutical composition comprising as an active ingredient a compound of claim 1 and a pharmaceutically acceptable carrier.

Claim 11 (previously canceled).

Claim 12 (previously canceled).

Claim 13 (currently amended): A method of treating airway and gastrointestinal disorders inflammation, allergy, nasal congestion, diseases of the GI-tract, cardiovascular disease, or disturbances of the central nervous system as well as allergy-induced airway responses, and obesity, said method comprising administering to a mammalian patient in need of such treatment a pharmaceutical composition which comprises therapeutically effective amounts of a compound of claim 1.

Claim 14 (previously canceled).

Claim 1/5 (currently Amended): A method of preparing a pharmaceutical composition for treating airway and gastrointestinal disorders inflammation, allergy, nasal congestion, diseases of the GI-tract, cardiovascular disease, or disturbances of the central nervous system as well as allergy-induced airway responses, and obesity, said method comprising bringing into intimate contact a compound of claim 1 and a pharmaceutically acceptable carrier. A compound exhibiting H₃ antagonist Claim 18 (currently amended):

activity, or enantiomers, stereoisomers and tautomers of said compound, or pharmaceutically acceptable salts or solvates of said compound, said compound being selected from the compounds with structures listed below:

Claim 17 (currently amended): A compound exhibiting both H₁ and H₃ antagonist activity, or enantiomers, stereoisomers and tautomers of said compound, or pharmaceutically acceptable salts or solvates of said compound, said compound being selected from the compounds with structures listed below:

Claim 18 (currently amended): A pharmaceutical composition for treating airway and gastrointestinal disorders inflammation, allergy, nasal congestion, diseases of the GI-tract, cardiovascular disease, or disturbances of the central nervous system as well as allergy induced airway responses, and obesity, said composition comprising therapeutically effective amount of a compound of claim 16 or claim 17 and a pharmaceutically acceptable carrier.

Claims 19-21 (canceled).

Claim 2/2 (new):

The compound of claim 1, wherein M is a piperidine.

Claim 23 (new):

The compound of claim 1, wherein M is a pyrrolidine.